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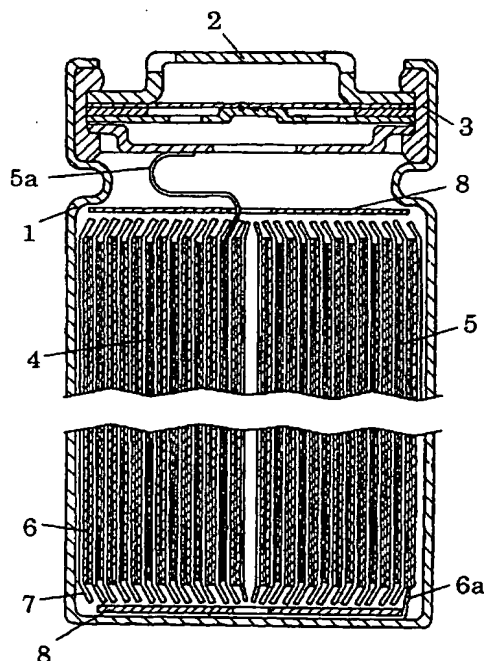
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(54) **NONAQUEOUS ELECTROLYTE SECONDARY CELL**

(57) A non-aqueous electrolyte secondary battery using composite particles for its negative electrode. In the composite particles, nucleus particles including at least one element selected from tin, silicon, and zinc as their constituent element are entirely or partly covered with a solid solution or inter-metallic compound of said constituent element and at least one element selected from groups consisting of Group 2 elements, transition elements, and Group 12, Group 13, and Group 14 elements in the Periodic Table except for the constituent element of the nucleus particles and carbon. Further, the present invention is characterized in that the NMR signals of the lithium intercalated in the composite particles appear within the range of -10 to 40 ppm with respect to lithium chloride and at least one signal appears within the range of -10 to 4 ppm

**FIG. 1**



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## Description

## FIELD OF THE INVENTION

5 [0001] The present invention relates to non-aqueous electrolyte secondary batteries, particularly to non-aqueous electrolyte secondary batteries (hereinafter referred to as "secondary batteries") with high energy density and improved electrochemical characteristics, such as charge/discharge capacity and cycle life, that are provided by the improved negative electrode material and non-aqueous electrolyte.

## 10 BACKGROUND OF THE INVENTION

[0002] High electromotive force and high energy density are featured by lithium secondary batteries recently used for such mobile communications equipment as personal digital assistants and mobile electronic apparatus, main power supply for the mobile electronic gears, small domestic power storage devices, and motor bicycles, electric vehicles, and hybrid cars using motors as their driving sources.

15 [0003] Lithium ion secondary batteries using organic electrolytic solutions, carbon materials as their negative electrode active materials, and lithium-containing composite oxides as their positive electrode active materials have higher energy density and more excellent low-temperature characteristics than secondary batteries using aqueous solutions. Moreover, without using lithium metals for their negative electrodes, lithium ion secondary batteries also have excellent cycle stability and safety, thus rapidly becoming commercially practical. Lithium polymer batteries using electrolytes like macromolecular (polymer) gel containing organic electrolytic solutions are also being developed as a new battery family of thin and light type.

20 [0004] When a lithium metal with high capacity is used as negative electrode material, dendrite-like deposition is formed on the negative electrode during charging. During repeated charge and discharge operations, the dendrite may penetrate the separator or polymer gel electrolyte, reach the positive electrode side and thus cause internal short circuits. Having a large specific surface area and thus high reactivity, the lithium deposition reacts with the solvent, becomes inactive, and decreases the charge/discharge efficiency. This phenomenon increases the internal resistance of the battery and produces particles isolated from electronic conduction network, thus decreasing the charge/discharge efficiency of the battery. For these reasons, lithium secondary batteries using lithium metals as their negative electrode materials have problems of having low reliability and poor cycle life characteristics.

30 [0005] At present, lithium secondary batteries use, for their negative electrodes, carbon materials capable of intercalating and de-intercalating lithium ions, and have become commercially available. Generally, since metal lithium does not deposit on a carbon negative electrode, it does not cause the problem of internal short circuits resulting from the production of dendrite. However, the theoretical capacity of graphite, one of carbon materials now in use, is 372 mAh/g, which is so small as one-tenth the theoretical capacity of pure Li metal.

35 [0006] As other negative electrode materials, metallic and non-metallic pure elements that form compounds with lithium are known. For example, the composition formula of a compound of tin (Sn), silicon (Si), or zinc (Zn) containing largest amount of lithium is expressed by  $\text{Li}_{22}\text{Sn}_5$ ,  $\text{Li}_{22}\text{Si}_5$ , or  $\text{LiZn}$ , respectively. Within this composition range, generally, no metallic lithium deposits; therefore, there is no problem of internal short circuits caused by the formation of dendrite. The electrochemical capacities between these compounds and each of the pure elements are 993 mAh/g, 4,199 mAh/g, and 410 mAh/g, respectively, all of which are larger than that of graphite.

40 [0007] Other compounds for negative electrode materials proposed include a nonferrous metal silicide consisting of transition elements disclosed in Japanese Patent Laid-Open Publication No. H07-240201, and a negative electrode material consisting of an inter-metallic compound containing at least one element selected from Group 4B elements, P, and Sb and having a crystal structure of one of  $\text{CaF}_2$ , ZnS, or  $\text{AlLiSi}$  type disclosed in Japanese Patent Laid-Open Publication No.H09-63651.

45 [0008] However, each of the aforementioned negative electrode materials with high capacity has the following problems.

50 [0009] Generally, the negative electrode materials of metallic and non-metallic pure elements that form compounds with lithium exhibit poorer charge/discharge cycle characteristics than carbon negative electrode materials. This is probably because the negative electrode materials break by their volumetric expansion and shrinkage.

55 [0010] Meanwhile, unlike the aforementioned pure elements, a negative electrode material that consists of a non-ferrous metal silicide consisting of transition elements and an inter-metallic compound that contains at least one element selected from Group 4B elements, P, and Sb and has a crystal structure of one of  $\text{CaF}_2$ , ZnS, or  $\text{AlLiSi}$  type are proposed as negative electrode materials with improved cycle life characteristics in Japanese Patent Laid-Open Publication No. H07-240201 and No.H09-63651, respectively. In Japanese Patent laid-Open Publication No.H10-208741, the range of nuclear magnetic resonance (hereinafter abbreviated as NMR) signals of lithium intercalated in the negative electrode material is proposed.

[0011] In the battery using the nonferrous metal silicide consisting of transition elements as the negative electrode material disclosed in Japanese Patent Laid-Open Publication No.H07-240201, while the discharge capacities at the first, fiftieth and hundredth cycles shown in its examples and a comparative example indicate that its charge/discharge cycle characteristics have been more improved than those of the lithium metal negative electrode material, the discharge capacities have only increased by about 12 % at maximum compared with that of the natural graphite negative electrode material.

[0012] For the material disclosed in Japanese Patent Laid-Open Publication No.H09-63651, its examples and comparative examples show that it has more improved charge/discharge cycle characteristics than the negative electrode material of Li-Pb alloy and has higher discharge capacity than graphite. However, the battery considerably decreases its discharge capacity at 10 to 20 charge/discharge cycles and even with  $Mg_2Sn$  that is considered best, its capacity decreases to 70 % of its initial capacity after about 20 cycles. For this reason, this material is inferior in charge/discharge characteristics.

[0013] For the material disclosed in Japanese Patent Laid-Open Publication No. H10-208741, NMR signals of the lithium intercalated in various alloys appear within the range of 5 to 40 ppm. By using this material, an electrode material with high energy density and excellent cycle life is proposed. However, as for the cycle life of the battery using this electrode material, its capacity decreases to 70 % of its maximum after 372 cycles even when  $LiCoO_2$  is used for its positive electrode.

[0014] The present invention addresses the aforementioned problems conventional batteries have.

## SUMMARY OF THE INVENTION

[0015] The negative electrode of the battery of the present invention is characterized by the use of composite particles. In the composite particles, nucleus particles containing at least one element selected from tin, silicon, and zinc as their primary constituent element are covered with a solid solution or inter-metallic compound of the elements constituting the nucleus particles and at least one element selected from groups consisting of Group 2 elements, transition elements, and Group 12, Group 13, and Group 14 elements in the Periodic Table except for carbon and constituent element of the nucleus particles. Further, the negative electrode of the battery of the present invention is characterized in that NMR signals of the lithium intercalated in the negative electrode material appear within the range of -10 to 40 ppm with respect to lithium chloride as a reference and at least one NMR signal appears within the range of -10 to 4 ppm. In addition, the negative electrode of the battery of the present invention is characterized in that the above NMR signals with respect to lithium chloride as a reference appear within the range of -10 to 4 ppm and -10 to 20 ppm and the NMR signal strength appearing within the range of -10 to 4 ppm is 1 to 10 times as strong as those appearing within the range of 10 to 20 ppm.

[0016] With the aforementioned structure, non-aqueous electrolyte secondary batteries that address the problems of conventional batteries and have higher energy density and more excellent cycle characteristics can be provided.

## BRIEF DESCRIPTION OF THE DRAWINGS

[0017] FIG.1 is a longitudinal section of a cylindrical battery of the present invention.

## PREFERRED EMBODIMENTS OF THE INVENTION

[0018] The battery of the present invention has positive and negative electrodes capable of intercalating and de-intercalating lithium, a non-aqueous electrolytic solution and a separator or a solid electrolyte.

[0019] The negative electrode material of the present invention uses composite particles in which nucleus particles comprising solid phase A are entirely or partly covered with solid phase B.

[0020] Solid phase A includes of at least one element selected from tin, silicon, and zinc.

[0021] Solid phase B includes a solid solution or an inter-metallic compound of at least one element selected from tin, silicon, and zinc and the other elements. The other elements constituting solid phase B comprise at least one element selected from groups consisting of Group 2 elements, transition elements, and Group 12, Group 13, and Group 14 elements in the periodic table except for the above constituent element and carbon.

[0022] In the following, the above negative electrode material of the present invention is referred to as "composite particles". By using the above composite particles as negative electrode material, solid phase B inhibits the expansion and shrinkage caused by charge and discharge of solid phase A, and thus the negative electrode material with excellent charge/discharge cycle characteristics can be obtained.

[0023] In the negative electrode material of the present invention, solid phase A is considered to mainly contribute to increase charge/discharge capacity because it contains at least one element selected from tin, silicon, and zinc that have high capacity. Solid phase B entirely or partly covering the nucleus particles consisting of solid phase A contributes

to the improvement in charge/discharge cycle characteristics. The amount of lithium contained in solid phase B is generally smaller than that contained in the pure metal, solid solution, or inter-metallic compound.

[0024] In other words, the negative electrode material used in the present invention is made of nucleus particles comprising at least one element selected from tin, silicon, and zinc that provide high capacity, and a solid solution or inter-metallic compound that is unlikely to pulverize and covers the nucleus particles. The solid solution or inter-metallic compound of the covering layer can restrain a change in the crystal structure of the nucleus particles, i.e. a large volumetric change, thus inhibiting the pulverization of the nucleus particles.

[0025] When the negative electrode materials react with lithium, lithium exists in the composite particles with covalent bonds or as ions. The NMR signals of covalent bonded  $^7\text{Li}$  appear within the range of 10 to 20 ppm. The site in which covalent bonded lithium exists is hereinafter referred to as "X site". The NMR signals of  $^7\text{Li}$  ions appear within the range of -10 and 40 ppm. The site in which lithium ion exists is hereinafter referred to as "Y site".

[0026] The Y site partly includes the signals related to irreversible factors. Such signals are generally found for negative electrodes of carbon materials, and seldom found for those of the present invention.

[0027] As described above, when a large amount of reversible X and Y sites of lithium exist, high capacity and excellent cycle characteristics can be obtained. In other words, appearance of the NMR signals of  $^7\text{Li}$  within the range of -10 to 40 ppm and -10 to 4 ppm at the same time in a charged state show the existence of both X and Y sites. It was found that high capacity and excellent cycle characteristics can be attained in such a state.

[0028] Moreover, it was found that higher capacity and more excellent cycle characteristics can be attained when the strength of signals appearing within the range of -10 to 4 ppm indicating the existence of the Y site are 1 to 10 times as strong as those of the signals appearing within the range of 10 to 20 ppm indicating the existence of the X site.

[0029] As described above, the present invention discriminates how lithium has been intercalated in the negative electrode material, i.e. as ions or covalent bonded, and determines the existence ratio using the NMR signals of  $^7\text{Li}$  to provide the material with high capacity and excellent cycle characteristics.

[0030] In other words, in order to exhibit high capacity and excellent cycle characteristics, the NMR signals of  $^7\text{Li}$  of the negative electrode material should appear within the range of -10 to 40 ppm and -10 to 4 ppm in a charged state.

[0031] The NMR signals of  $^7\text{Li}$  appearing within the range of -10 to 40 ppm shows the existence of  $^7\text{Li}$  ions and covalent bonded  $^7\text{Li}$  mixed. The NMR signals of  $^7\text{Li}$  appearing within the range of -10 and 4 ppm shows the existence of lithium ions that provides high capacity. Moreover, when the signals appearing within the range of -10 to 4 ppm that indicates the existence of lithium ions are 1 to 10 times as strong as those appearing within the range of 10 to 20 ppm that indicates the existence of covalent bonded lithium, much higher capacity can be attained.

[0032] The positive and negative electrodes used for the present invention are made by applying the surface of a current collector with a layer of an electrode mixture. The mixture contains positive electrode active materials or the above negative electrode materials capable of electrochemically and reversibly intercalating and de-intercalating lithium ions together with a conductive material, binder, and the like.

[0033] Next, producing methods of composite particles used as the negative electrode materials are described.

[0034] One method of producing composite particles includes:

quenching and solidifying the melt containing each element constituting the composite particles in the composition ratios, using dry atomization, wet atomization, roll quenching, rotating electrode method, and other methods; and heat-treating the solidified material at a temperature lower than the solid line temperature of the solid solution or inter-metallic compound. The solid line temperature is determined by the composition ratio.

[0035] Quenching and solidifying the melt allows particles of solid phase A to deposit as the nucleus of the particles and solid phase B to deposit so as to entirely or partly cover the particles of solid phase A. Thereafter, heat treatment can improve the homogeneity of each of the solid phases A and B. Even when the above heat treatment is not performed, composite particles appropriate for the present invention can be obtained. Another cooling method can be used on condition that it can sufficiently cool the melt at a high speed.

[0036] Another production method is forming a layer comprising elements required for forming solid phase B over the surfaces of the powder of solid phase A, and heat-treating it at a temperature lower than the solid line temperature. This heat treatment allows component elements in solid phase A to diffuse toward the layer and to form solid phase B as a covering layer. The layer is formed using plating, mechanical alloying, or other method. The mechanical alloying method does not require heat treatment. Any method capable of forming the layer can be used.

[0037] Conductive materials for the negative electrodes can be any electronic conductive materials. Examples of such materials include: graphite materials including natural graphite (scale-like graphite), artificial graphite, and expanded graphite; carbon black materials such as acetylene black, high-structure carbon black typified by Ketjen black, channel black, furnace black, lamp black and thermal black; conductive fibers such as carbon fibers and metal fibers; metal powders such as copper and nickel; and organic conductive materials such as polyphenylene derivatives. These materials can be used independently or in combination. Among these conductive materials, artificial graphite, acetylene

black, and carbon fibers are especially preferable.

[0038] The amount of conductive materials to be added is not specifically defined; however, 1 to 50 wt %, especially 1 to 30 wt %, of the negative electrode material is preferable. Having electronic conductivity in itself, the negative electrode material of the present invention (composite particles) can work as a battery electrode material without any additional conductive materials. Thus, more composite particles can be included in the electrode by the amount of conductive materials.

[0039] For binders for the negative electrodes, both thermoplastic resin and thermosetting resin can be used. Preferable binders for the present invention include the following materials: polyethylene, polypropylene, polytetrafluoroethylene (PTFE), polyvinylidene fluoride (PVDF), styrene-butadiene rubber, a tetrafluoroethylene-hexafluoropropylene copolymer (FEP), a tetrafluoroethylene-perfluoroalkyl vinyl ether copolymer (PFA), a vinylidene fluoride-hexafluoropropylene copolymer, a vinylidene fluoride-chlorotrifluoroethylene copolymer, an ethylene-tetrafluoroethylene copolymer (ETFE resin), polychlorotrifluoroethylene (PCTFE), a vinylidene fluoride-pentafluoropropylene copolymer, a propylene-tetrafluoroethylene copolymer, an ethylene-chlorotrifluoroethylene copolymer (ECTFE), a vinylidene fluoride-hexafluoropropylene-tetrafluoroethylene copolymer, a vinylidene fluoride-perfluoromethyl vinyl ether-tetrafluoroethylene copolymer, an ethylene-acrylic acid copolymer or its Na<sup>+</sup> ion crosslinking body, an ethylene-methacrylic acid copolymer or its Na<sup>+</sup> ion crosslinking body, an ethylene-methyl acrylate copolymer or its Na<sup>+</sup> ion crosslinking body, and an ethylene-methyl methacrylate copolymer or its Na<sup>+</sup> ion crosslinking body. These materials can be used independently or in combination. Preferable materials among these materials are styrene-butadiene rubber, polyvinylidene fluoride, an ethylene-acrylic acid copolymer or its Na<sup>+</sup> ion crosslinking body, an ethylene-methacrylic acid copolymer or its Na<sup>+</sup> ion crosslinking body, an ethylene-methyl acrylate copolymer or its Na<sup>+</sup> ion crosslinking body, and an ethylene-methyl methacrylate copolymer or its Na<sup>+</sup> ion crosslinking body.

[0040] For current collectors for the negative electrodes, any electronic conductive material can be used on condition that it does not chemically change in the battery constructed. Examples of such materials include stainless steel, nickel, copper, titanium, carbon, and conductive resin as well as copper or stainless steel surface-treated with carbon, nickel, or titanium. Among these, copper and copper alloys are especially preferable. These materials can also be used after their surfaces have been oxidized. Desirably, the current collector is surface-treated to have a roughened surface. The shapes of the current collector include a film, sheet, net, punched sheet, lath, porous body, foam, and formed and grouped fibers as well as a foil. The thickness is not specifically defined; however, collectors 1 to 500  $\mu\text{m}$  thick are used.

[0041] As positive electrode active materials, compounds containing lithium or containing no lithium can be used. Such compounds include  $\text{Li}_x\text{CoO}_2$ ,  $\text{Li}_x\text{NiO}_2$ ,  $\text{Li}_x\text{MnO}_2$ ,  $\text{Li}_x\text{Co}_y\text{Ni}_{1-y}\text{O}_2$ ,  $\text{Li}_x\text{Co}_y\text{M}_{1-y}\text{O}_2$ ,  $\text{Li}_x\text{Ni}_{1-y}\text{M}_y\text{O}_2$ ,  $\text{Li}_x\text{Mn}_2\text{O}_4$ , and  $\text{Li}_x\text{Mn}_{2-y}\text{M}_y\text{O}_4$  (M is at least one of Na, Mg, Sc, Y, Mn, Fe, Co, Ni, Cu, Zn, Al, Cr, Pb, Sb and B, where  $x = 0-1.2$ ,  $Y = 0-0.9$ ,  $z = 2.0-2.3$ ). The above value  $x$  is the value before charging and discharging; thus it changes along with charging and discharging. Other usable positive electrode materials include transition metal chalcogenides, a vanadium oxide and its lithium compounds, a niobium oxide and its lithium compounds, a conjugate polymer using organic conductive materials, and shevrit phase compounds. It is also possible to use a plurality of different positive electrode materials in combination. The average diameter of particles of the positive electrode active materials is not specifically defined; however, desirably it is 1-30  $\mu\text{m}$ .

[0042] Conductive materials for the positive electrode can be any electronic conductive material on condition that it does not chemically change within the range of charge and discharge electric potentials of the positive electrode materials in use. Examples of such materials include: graphite materials including natural graphite (scale-like graphite) and artificial graphite; carbon black materials such as acetylene black, high-structure carbon black typified by Ketjen black, channel black, furnace black, lamp black and thermal black; conductive fibers such as carbon fibers and metal fibers; metal powders such as fluorinated carbon and aluminum; conductive whiskers such as a zinc oxide and potassium titanate; conductive metal oxides such as a titanium oxide; and organic conductive materials such as polyphe-nylene derivatives. These materials can be used independently or in combination. Among these conductive materials, artificial graphite and acetylene black are especially preferable. The amount of conductive materials to be added is not specifically defined; however, 1 to 50 wt %, especially 1 to 30 wt %, of the positive electrode material is preferable. For carbon and graphite, 2 to 15 wt % is especially preferable.

[0043] For binders for the positive electrodes, both thermoplastic and thermosetting resins can be used. The aforementioned binders for the negative electrodes can preferably be used, and more preferable materials are PVDF and PTFE.

[0044] For current collectors for the positive electrodes, any electronic conductive material can be used on condition that it does not chemically change within electric potentials in the range of charge and discharge of the positive electrode material in use. For example, the aforementioned current collectors for the negative electrode can preferably be used. The thickness is not specifically defined; however, collectors 1 to 500  $\mu\text{m}$  thick are used.

[0045] The electrode mixtures for the negative and positive electrode plates, can include filler, dispersant, ion conductor, pressure intensifier and other various additives as well as a conductive material and binder. As the filler, any fibrous material that does not chemically change in the constructed battery can be used. Generally, olefinic polymers

such as polypropylene and polyethylene and such fibers as glass fibers and carbon fibers are used. The amount of the filler to be added is not specifically defined; however, 0 to 30 wt % of the electrode mixture is preferable.

[0046] Preferably, the negative and positive electrodes are structured so that at least the surface made of the negative electrode mixture is opposite the surface made of the positive electrode mixture.

[0047] The non-aqueous electrolytes used for the present invention are composed of non-aqueous solvents and lithium salts dissolved in the solvents. Examples of such non-aqueous solvents include: cyclic carbonates such as ethylene carbonate (EC), propylene carbonate (PC), butylene carbonate (BC), and vinylene carbonate (VC); acyclic carbonates such as dimethyl carbonate (DMC), diethyl carbonate (DEC), ethylmethyl carbonate (EMC), and dipropyl carbonate (DPC); aliphatic carboxylates such as methyl formate, methyl acetate, methyl propionate, and ethyl propionate;  $\gamma$ -lactones such as  $\gamma$ -butyrolactone; acyclic ethers such as 1,2 - dimethoxy ethane (DME), 1,2 - diethoxy ethane (DEE), and ethoxy methoxy ethane (EME); cyclic ethers such as tetrahydrofuran and 2 - methyl tetrahydrofuran; and non-protonic organic solvents such as dimethyl sulfoxide, 1,3 - dioxolane, formamide, acetamide, dimethylformamide, dioxolane, acetonitrile, propyl nitrile, nitromethane, ethyl monoglime, triester of phosphoric acid, trimethoxy methane, dioxolane derivatives, sulfolane, methylsulfolane, 1,3-dimethyl-2-imidazolidinone, 3-methyl-2-oxazolidinone, propylene carbonate derivatives, tetrahydrofuran derivatives, ethyl ether, 1,3-propane saltane, anisole, dimethyl sulfoxide and N-methylpyrrolidone. These solvents are used independently or as a mixture of one, two or more solvents. Among these, mixtures of cyclic carbonate and acyclic carbonate, or mixtures of cyclic carbonate, acyclic carbonate and aliphatic carboxylate are especially preferable.

[0048] Lithium salts dissolved into the above solvents include  $\text{LiClO}_4$ ,  $\text{LiBF}_4$ ,  $\text{LiPF}_6$ ,  $\text{LiAlCl}_4$ ,  $\text{LiSbF}_6$ ,  $\text{LiSCN}$ ,  $\text{LiCl}$ ,  $\text{LiCF}_3\text{SO}_3$ ,  $\text{LiCF}_3\text{CO}_2$ ,  $\text{Li}(\text{CF}_3\text{SO}_2)_2$ ,  $\text{LiAsF}_6$ ,  $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ ,  $\text{LiB}_{10}\text{Cl}_{10}$ , lithium salt of lower aliphatic carboxylic acid,  $\text{LiCl}$ ,  $\text{LiBr}$ ,  $\text{LiI}$ , chloroboron lithium, 4-phenyl boric acid, and an imide group. These lithium salts can be dissolved in the above non-aqueous solvents individually or in combination of two or more lithium salts to use as an electrolytic solution. It is especially preferable to include  $\text{LiPF}_6$  in the electrolytic solution.

[0049] Especially preferable non-aqueous electrolytes of the present invention includes at least EC and EMC, and as a supporting salt,  $\text{LiPF}_6$ . The amount of the electrolyte to be added to the battery is not specifically defined. Considering the amount of the positive and negative electrode materials and the size of the battery, the required amount is used. The amount of the supporting electrolytes against the non-aqueous solvents is not specifically defined; however, 0.2 to 2 mol/l is preferable, and especially 0.5 to 1.5 mol/l is more preferable.

[0050] Besides the electrolytic solutions, the following solid electrolytes can also be used. The solid electrolytes are classified into inorganic and organic ones.

[0051] Well known inorganic solid electrolytes are lithium nitrides, lithium halides, and lithium oxides, and the like. Among them,  $\text{Li}_4\text{SiO}_4$ ,  $\text{Li}_4\text{SiO}_4\text{-LiI-LiOH}$ ,  $\text{xLi}_3\text{PO}_4\text{-(1-x)Li}_4\text{SiO}_4$ ,  $\text{Li}_2\text{SiS}_3$ ,  $\text{Li}_3\text{PO}_4\text{-Li}_2\text{S-SiS}_2$  and phosphorus sulfide compounds can be used effectively.

[0052] As the organic solid electrolytes, polymer materials such as polyethylene oxides, polypropylene oxides, polyphosphazene, polyaziridine, polyethylene sulfides, polyvinyl alcohol, polyvinylidene fluorides, polyhexafluoropropylene and their derivatives, mixtures and complexes can effectively be used.

[0053] Moreover, it is effective to add other compounds to the electrolytic solution in order to improve discharge and charge/discharge characteristics. Such compounds include triethyl phosphite, triethanolamine, cyclic ethers, ethylene diamine, n-glime, pyridine, triamide hexaphosphate, nitrobenzene derivatives, crown ethers, quaternary ammonium salts, and ethylene glycol dialkyl ether.

[0054] As separators, insulating micro-porous thin film having large ion permeability and predetermined mechanical strength is used. Desirably, these separators have functions of closing their pores and increasing resistance above predetermined temperatures. In terms of resistance to organic solvents and hydrophobic properties, olefinic polymers such as polypropylene and polyethylene are used independently or in combination, or sheets and woven or non-woven fabrics made of glass fibers, and the like, are used. Desirably, the micro-porous separator has pores of a diameter that does not allow the positive or negative electrode material, binder, and conductive material liberated from the electrode sheets to pass. For example, pores 0.01 to 1  $\mu\text{m}$  in diameter are preferable. Generally, separators 10 to 300  $\mu\text{m}$  thick are used. The porosity of the separator is determined according to the permeability of electrons and ions, materials, and thickness. Generally, a porosity of 30 to 80 % is desirable.

[0055] A battery with integrated porous separator and positive and negative electrodes can be constructed. The integrated positive and negative electrode mixtures contain a polymer material that absorbs and holds an organic electrolytic solution composed of a solvent and a lithium salt dissolved in the solvent. The separator is composed of a polymer absorbing and holding the organic electrolytic solution. In this case, any polymer material capable of absorbing and holding the organic electrolytic solution can be used. Especially, a vinylidene fluoride - hexafluoropropylene copolymer is preferable.

[0056] The battery can be of any shape including a coin, button, sheet, lamination, cylinder, flat, square, and a large type used for an electric vehicle.

[0057] The materials used for the batteries of the present invention are hereinafter described in detail. However, the

present invention is not limited to these examples.

[0058] The positive and negative electrodes used for the batteries of the present invention are made of current collectors coated with the mixtures. The mixtures essentially include positive electrode active material and negative electrode material capable of electrochemically and reversibly intercalating and de-intercalating lithium ions, and conductive materials and binder.

#### Production of composite particles

[0059] Table 1 shows the composition of solid phases A and B of the composite particles (single element, inter-metallic compound, and solid solution) used for the examples of the present invention, ratios of the elements at preparation, fusing temperatures, and solid line temperatures.

[0060] Powder or a block of each element constituting composite particles was put into a crucible in the composition ratio shown in Table 1. The material was fused at a temperature shown in Table 1 and quenched using the roll quenching method and solidified to obtain a solidified material. Sequentially, the solidified material was heat-treated in an inert atmosphere at a temperature about 10 to 50 °C lower than the solid line temperature shown in Table 1 for 20 hours. The heat-treated material was pulverized with a ball mill and classified with a sieve to obtain composite particles 45 µm or smaller in size. Electron microscope observations verified that these particles were made of particles of solid phase A entirely or partly covered with particles of solid phase B.

[0061] The structure of the batteries of the present invention is described with reference to specific examples of a cylindrical battery.

#### Example 1

[0062] FIG.1 shows a longitudinal section of the cylindrical battery of the present invention. Positive electrode plate 5 and negative electrode plate 6 are spirally wound via separator 7 and housed in battery case 1. Positive electrode lead 5a is led out from the positive electrode plate 5 and connected to sealing plate 2. Negative electrode lead 6a is led out from negative electrode plate 6 and connected to the bottom of battery case 1.

[0063] Metals and alloys having electronic conductivity and resistance to organic electrolytic solutions can be used for the battery case and lead plate. For example, metals such as iron, nickel, titanium, chromium, molybdenum, copper, and aluminum, and their alloys are used. Stainless steel plates or processed Al-Mn alloy plates are especially preferable materials for the battery case, aluminum for the positive electrode lead, and nickel for the negative electrode lead. For the battery case, various kinds of engineering plastics and those in combination with metals can be used to reduce its weight.

[0064] Insulating rings 8 are provided on the top and bottom of grouped electrode plates 4, respectively. The sealing plate can be used as a safety valve. Conventionally known various safety elements can be installed other than the safety valve. Such safety elements include a fuse, bimetal, and PTC device functioning as an overcurrent protection device. Other than the safety valve, notching the battery case, cracking the gasket, cracking the sealing plate, or breaking connection with the lead can also be used to prevent an increase in internal pressure of the battery case. A protection circuit capable of protecting from overcharge or overdischarge can be incorporated in a battery charger or provided independently. In addition, a system in which an increase in the internal pressure of the battery case breaks the current can be incorporated to eliminate overcharge. In this case, compounds capable of increasing the internal pressure can be contained in the electrode mixtures or electrolytes. Such compounds include carbonates such as  $\text{Li}_2\text{CO}_3$ ,  $\text{LiHCO}_3$ ,  $\text{Na}_2\text{CO}_3$ ,  $\text{NaHCO}_3$ ,  $\text{CaCO}_3$ , and  $\text{MgCO}_3$ .

[0065] For welding the cap, battery case, sheet and lead, well-known methods (e.g. electric welding using direct or alternating current, laser welding, and ultrasonic welding) can be used. As the sealing materials for sealing the openings, conventionally known compounds and mixtures, such as asphalt, can be used.

[0066] Negative electrode plate 6 was made by:

mixing 20 wt % of carbon powder, 5 wt % of PVDF and 75 wt% of composite particles synthesized under the above conditions;

dispersing the mixture in dehydrated N-methyl pyrrolidinone to prepare a slurry;

coating the negative electrode current collector made of a copper foil with the slurry; and  
drying and then rolling it.

[0067] Positive electrode plate 5 was made by:

mixing 10 wt % of carbon powder, 5 wt % of PVDF and 85 wt % of lithium cobaltate powder;

dispersing the mixture in dehydrated N-methylpyrrolidinone to prepare a slurry;

coating the positive electrode current collector made of a copper foil with the slurry; and drying and then rolling it.

[0068] An electrolytic solution containing 1.5 mol/l of  $\text{LiPF}_6$  dissolved in a mixed solvent of EC and EMC in a volumetric ratio of EC:EMC = 1:1 was used.

[0069] Thus, the batteries were made using the materials shown in Table 1 for their negative electrodes. The cylindrical batteries produced are 18 mm in diameter and 65 mm in height.

#### Comparative Example 1

[0070] A battery using graphite for its negative electrode was produced in the same manner as Example 1 for comparison.

[0071] After these batteries were charged at a constant current of 100 mA, they were discharged at a constant current of 100 mA to 2.0 V. Such charge/discharge cycles were repeated in an oven at a constant temperature of 20 °C. The charge/discharge cycles were repeated up to 400 times and the ratio of the discharge capacity at the 400th cycle to the initial discharge capacity was shown as a capacity retention rate in Table 2.

[0072] In addition, NMR measurement was performed on the  $^7\text{Li}$  of the negative electrode materials in a charged state and the NMR signals of the intercalated lithium were measured with respect to the lithium chloride as a reference. Several NMR signals appeared within the range of -10 to 40 ppm. FIG. 2 shows the positions at which the signals appeared and the signal strength ratio of the NMR signals appearing within the range of -10 to 4 ppm to those appearing within the range of 10 to 20 ppm. Due to variations from lot to lot, the same material exhibited different signal positions and intensities. The NMR measuring conditions of the  $^7\text{Li}$  are as follows:

Instrument: INOVA 400 (VARIAN Co.)

Temperature: room temperature

Measured nuclear:  $^7\text{Li}$

Revolution of the samples: 10 kHz

Reference Material:  $\text{LiCl}$  1M

[0073] Table 2 shows the followings about batteries A to S using materials A to S as negative electrode materials.

[0074] Even when the NMR signals of  $^7\text{Li}$  intercalated in the negative electrode materials appear within the range of -10 to 40 ppm,

(1) in the case where no signals appear within the range of -10 to 4 ppm and

(2) in the case where all the signals appear in the magnetic field lower than 4 ppm,

the capacity retention rates after 400 cycles are high; however, the initial capacities and the capacities after 400 cycles are lower than those of battery T using the graphite negative electrode.

[0075] In the case where the signals appear between the magnetic fields lower than 5 ppm and higher than -10 ppm, the initial capacities are large but the capacity retention rates after 400 cycles are small and the discharge capacities after 400 cycles are smaller than that of battery T.

[0076] Meanwhile, when the signals appear within the range of -10 to 40 ppm and at least one signal appears within the range of -10 to 4 ppm, the initial capacities are larger than that of battery T and the capacity retention rates after 400 cycles are 70 % or higher and the capacities after the cycles are larger than that of battery T.

[0077] When the signal strength of the signals appearing within the range of -10 to 4 ppm indicating the  $^7\text{Li}$  mainly in the ionic state is 1 to 10 times as strong as the signal strength of the signals appearing within the range of 10 to 20 ppm indicating the  $^7\text{Li}$  mainly covalent bonded, the capacities after 400 cycles are the largest.

[0078] As mentioned above, when the NMR signals of lithium intercalated in the negative electrode materials appear within the range of -10 to 40 ppm with respect to lithium chloride as a reference and at least one NMR signal appears within the range of -10 to 4 ppm, a secondary battery with high capacity and excellent cycle characteristics can be obtained. Moreover, when the NMR signals of the lithium intercalated in the negative electrode materials appearing within the range of -10 to 4 ppm are 1 to 10 times as strong as those appearing within the range of 10 to 20 ppm, a secondary battery with higher capacity and more excellent cycle characteristics can be obtained.

[0079] Regarding the elements constituting negative electrode materials, when solid phase A is made of tin, Mg selected from Group 2 elements, Fe and Mo from transition elements, Zn and Cd from Group 12 elements, In from Group 13 elements, and Pb from Group 14 elements are used. However, using other elements in each Group could provide similar results. Solid phase A may include any element other than tin, including a trace amount of such elements as O, C, N, S, Ca, Mg, Al, Fe, W, V, Ti, Cu, Cr, Co, and P.



[0080] Regarding the elements constituting negative electrode materials, when solid phase A is made of silicon, Mg selected from Group 2 elements, Co and Ni from transition elements, Zn from Group 12 elements, Al from Group 13 elements, and Sn from Group 14 elements are used. However, using other elements in each Group can provide similar results.

5 [0081] In addition, when solid phase A is made of zinc, Mg selected from Group 2 elements, Cu and V from transition elements, Cd from Group 12 elements, Al from Group 13 elements, and Ge from Group 14 elements are used. However, using other elements in each Group can provide similar results.

10 [0082] The composition ratios of the elements constituting the negative electrode materials are not specifically defined. Any composition can be used on condition that the elements provide two phases so that one phase (solid phase A) essentially consists of tin, silicon, and zinc, and the other phase (solid phase B) entirely or partly covers the surface of solid phase A. Solid phase A may contain any element except each constituent element, for example, a trace amount of such elements as O, C, N, S, Ca, Mg, Al, Fe, W, V, Ti, Cu, Cr, Co, and P.

15 [0083] As described above, a non-aqueous electrolyte secondary battery having higher capacity and more excellent cycle characteristics than conventional ones using carbon materials for their negative electrodes can be obtained, when the battery uses aforementioned composite particles composed of nucleus particles of solid phase A and solid phase B entirely or partly covering the solid phase A, and NMR signals of the lithium intercalated in the negative electrode material appear within the range of -10 to 40 ppm with respect to lithium chloride as a reference and at least one signal appears within the range of -10 to 4 ppm. Moreover, a non-aqueous electrolyte secondary battery with higher capacity and more excellent cycle characteristics than conventional ones using carbon materials for their negative electrodes can be obtained when the NMR signals of the lithium intercalated in the negative electrode material appearing within the range of -10 to 4 ppm with respect to lithium chloride as a reference is 1 to 10 times as strong as those appearing within the range of 10 to 20 ppm.

#### INDUSTRIAL APPLICABILITY

25 [0084] The secondary battery using the composite particles for the negative electrode in accordance with the present invention has higher energy density and more effects on the improvement in charge/discharge cycle life characteristics than the conventional batteries using carbon materials for their negative electrodes. For this reason, the secondary battery of the present invention can be used for personal digital assistants, mobile electronic equipment, small domestic power storage devices, motor bicycles, electric vehicles, hybrid cars, and the like, thus providing great industrial effects. The uses of the secondary battery of the present invention are not limited to those described above, and can be used for any application requiring a secondary battery.

Table 1

Negative electrode material	Solid phase A	Solid phase B	Fusing temperature(°C)	Solid line temperature(°C)	Composition ratio(atomic %)
Material A	Sn	Mg <sub>2</sub> Sn	770	204	Sn:Mg = 50:50
Material B	Sn	FeSn <sub>2</sub>	1540	513	Sn:Fe = 70:30
Material C	Sn	MoSn <sub>2</sub>	1200	800	Sn:Mo = 70:30
Material D	Sn	Solid solution of Zn and Sn	420	199	Sn:Zn = 90:10
Material E	Sn	Solid solution of Cd and Sn	232	133	Sn:Cd = 95: 5
Material F	Sn	Solid solution of In and Sn	235	224	Sn:In = 98: 2
Material G	Sn	Solid solution of Sn and Pb	232	183	Sn:Pb = 80: 20
Material H	Si	Mg <sub>2</sub> Si	1415	946	Si:Mg = 70: 30
Material I	Si	CoSi <sub>2</sub>	1495	1256	Si:Co = 85: 15
Material J	Si	NiSi <sub>2</sub>	1415	993	Si:Ni = 69: 31
Material K	Si	Solid solution of Si and Zn	1415	420	Si:Zn = 50: 50
Material L	Si	Solid solution of Si and Al	1415	577	Si:Al = 40: 60
Material M	Si	Solid solution of Si and Sn	1415	232	Si:Sn = 50: 50

Material N	Zn	Mg <sub>2</sub> Zn <sub>11</sub>	650	364	Zn:Mg = 92.9:7.8
Material O	Zn	Solid solution of Zn and Cu	1085	425	Zn:Cu = 97: 3
Material P	Zn	VZn <sub>11</sub>	700	420	Zn: V = 94: 6
Material Q	Zn	Solid solution of Zn and Cd	420	266	Zn:Cd = 50: 50
Material R	Zn	Solid solution of Zn and Al	661	381	Zn:Al = 90: 10
Material S	Zn	Solid solution of Zn and Ge	938	394	Zn:Ge = 97: 3

Table 2

Battery	Electrode material	Charging voltage (V)	NMR signal positions (ppm)	Strength ratio* (times)	Initial discharge capacity (mAh)	Discharge capacity after 400th cycle (mAh)	Capacity retention rate (%)
A	Material A	4.0	8,13	0.5	1081	864	80
		4.05	4,13	0.8	1332	1038	78
		4.1	-4,13	1	1566	1205	77
		4.15	-6,13	10	1633	1208	74
		4.18	-10,13	12	1659	1161	70
		4.2	-12,13	15	1766	512	29
B	Material B	4.0	7,13	0.4	1077	872	81
		4.05	4,13	0.9	1473	1148	78
		4.1	-2,13	1	1789	1359	76
		4.15	-5,13	10	1849	1386	75
		4.18	-10,13	12	1855	1298	70
		4.2	-13,13	15	1898	284	15
C	Material C	4.0	6,13	0.5	1067	853	80
		4.05	3,13	0.7	1600	1248	78
		4.1	-5,13	2	1825	1350	74
		4.15	-6,13	10	1834	1338	73
		4.18	-10,13	12	1847	1292	70
		4.2	-11,13	14	1859	390	21

D	Material D	4.0	6,13	0.5	1047	827	79
		4.05	4,13	0.8	1578	1230	78
		4.1	-4,13	1	1829	1371	75
		4.15	-4,13	8	1839	1360	74
		4.18	-10,13	12	1852	1296	70
		4.2	-12,13	15	1877	356	19
E	Material E	4.0	8,13	0.5	1001	800	80
		4.05	4,13	0.8	1613	1258	78
		4.1	-4,13	1	1861	1414	76
		4.15	-5,13	10	1874	1386	74
		4.18	-10,13	11	1876	1313	70
		4.2	-11,13	12	1899	338	21
F	Material F	4.0	8,13	0.5	1073	879	82
		4.05	4,13	0.8	1623	1265	78
		4.1	-4,13	1	1846	1420	77
		4.15	-6,13	10	1852	1389	75
		4.18	-9,13	11	1861	1302	70
		4.2	-12,13	12	1888	339	18
G	Material G	4.0	6,13	0.5	1054	832	79
		4.05	4,13	0.8	1572	1227	78
		4.1	-3,13	1.5	1833	1411	77
		4.15	-4,13	10	1848	1404	76
		4.18	-10,13	12	1855	1317	71
		4.2	-11,13	15	1899	474	25
H	Material H	4.0	6,13	0.5	1091	872	80

I	Material I	4.05	3,13	0.7	1702	1327	78
		4.1	-4,13	1	1931	1467	76
		4.15	-5,13	8	1945	1458	75
		4.18	-10,13	13	1958	1370	70
		4.2	-12,13	15	1968	413	21
		4.0	8,13	0.5	1059	857	81
J	Material J	4.05	4,13	0.8	1677	1308	78
		4.1	-4,13	2.1	1911	1433	75
		4.15	-6,13	10	1924	1404	73
		4.18	-10,13	12	1935	1354	70
		4.2	-12,13	15	1940	465	24
		4.0	7,13	0.5	1069	855	80
K	Material K	4.05	3,13	0.65	1546	1221	79
		4.1	-4,13	1	1926	1502	78
		4.15	-5,13	10	1973	1499	76
		4.18	-10,13	12	1974	1381	70
		4.2	-12,13	15	1985	476	24
		4.0	7,13	0.5	1069	855	80
L	Material L	4.05	3,13	0.8	1578	1230	78
		4.1	-4,13	1	1918	1476	77
		4.15	-4,13	10	1932	1449	75
		4.18	-10,13	12	1944	1360	70
		4.2	-12,13	13	1969	433	22
		4.0	8,13	0.5	1066	863	81
		4.05	4,13	0.8	1769	1379	78

M	Material M	4.1	-4,13	1	1921	1459	76
		4.15	-5,13	8	1943	1457	75
		4.18	-9,13	11	1966	1395	71
		4.2	-11,13	15	1989	576	29
		4.0	8,13	0.5	1021	816	80
		4.05	2,13	0.7	1682	1328	79
		4.1	-3,13	1	1902	1483	78
		4.15	-6,13	10	1923	1461	76
		4.18	-10,13	11	1945	1361	70
		4.2	-12,13	15	1981	455	23
N	Material N	4.0	8,13	0.5	1039	851	82
		4.05	4,13	0.8	1508	1176	78
		4.1	-4,13	1	1764	1358	77
		4.15	-5,13	9	1789	1341	75
		4.18	-10,13	12	1809	1266	70
		4.2	-12,13	13	1817	363	20
		4.0	8,13	0.5	1039	808	80
		4.05	4,13	0.8	1510	1177	78
O	Material O	4.1	-4,13	1	1751	1348	77
		4.15	-6,13	10	1810	1357	75
		4.18	-10,13	11	1815	1270	70
		4.2	-12,13	12	1828	530	29
		4.0	8,13	0.5	1002	801	80
		4.05	3,13	0.8	1474	1149	78
P	Material P	4.1	-4,13	1.5	1692	1302	77

Q	Material Q	4.15	-7,13	10	1755	1316	75
		4.18	-10,13	12	1799	1259	70
		4.2	-12,13	15	1805	415	23
		4.0	6,13	0.5	1099	868	79
		4.05	4,13	0.8	1508	1176	78
		4.1	-4,13	1	1719	1323	77
		4.15	-5,13	10	1755	1333	76
		4.18	-10,13	12	1800	1260	70
R	Material R	4.2	-12,13	15	1805	541	30
		4.0	8,13	0.5	1092	862	79
		4.05	3,13	0.8	1445	1141	78
		4.1	-4,13	1.2	1754	1350	77
		4.15	-6,13	10	1800	1368	76
		4.18	-10,13	12	1805	1281	71
		4.2	-12,13	14	1810	325	18
		4.0	8,13	0.5	1098	878	80
S	Material S	4.05	4,13	0.7	1455	1134	78
		4.1	-4,13	1	1710	1316	77
		4.15	-8,13	10	1745	1308	75
		4.18	-10,13	12	1795	1256	70
		4.2	-12,13	15	1809	434	24
T	Graphite	4.2	—	—	1510	906	60



## Reference Numerals

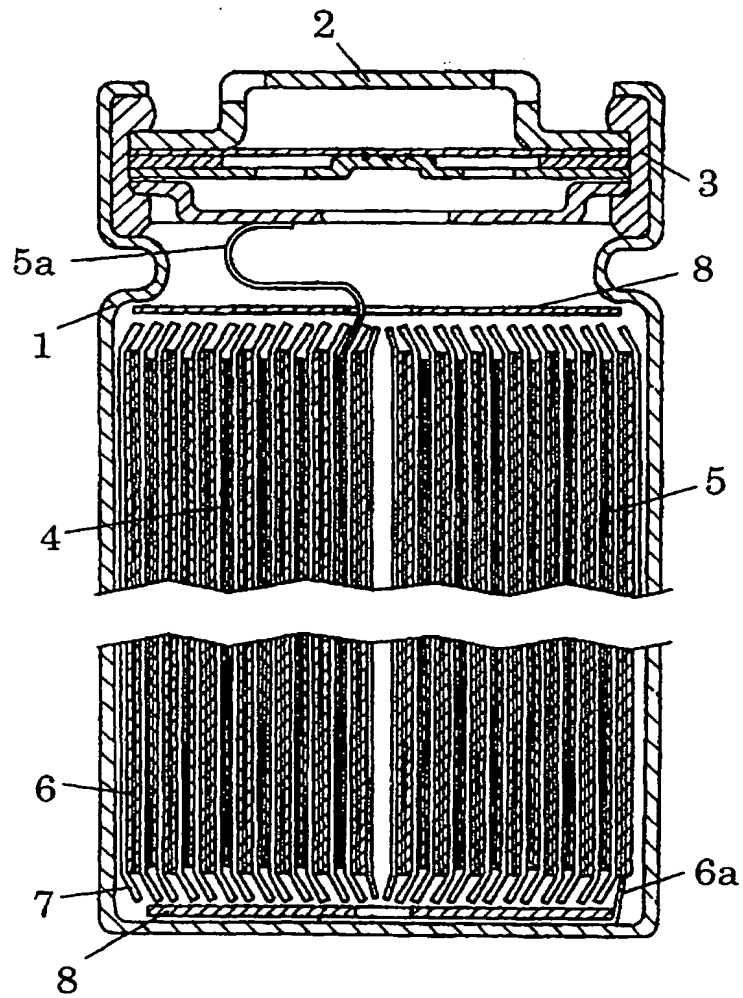
[0085]

- 5 1. battery case
- 2. sealing plate
- 3. insulating gasket
- 4. electrode plate group
- 5. positive electrode plate
- 10 5a. positive electrode lead
- 6. negative electrode
- 6a. negative electrode lead
- 7. separator
- 8. insulating ring

## Claims

- 20 1. A non-aqueous electrolyte secondary battery having a non-aqueous electrolyte , separator, and positive and negative electrodes capable of intercalating and de-intercalating lithium, said negative electrode including composite particles including at least one element selected from tin, silicon, or zinc as a constituent element thereof and at least one of:
  - 25 (a) a solid solution of said constituent element and at least one element selected from groups consisting of Group 2 elements, transition elements, Group 12, Group 13, and Group 14 elements in a Periodic Table except for said constituent element and carbon; and
  - (b) an inter-metallic compound of said constituent element and at least one element selected from groups consisting of Group 2 elements, transition elements, Group 12, Group 13, and Group 14 elements in the Periodic Table except for said constituent element and carbon,
  - 30 covering said particles entirely or partly,
  - wherein NMR signals of lithium intercalated in said composite particles appear within the range of -10 to 40 ppm with respect to lithium chloride as a reference and at least one signal appears within the range of -10 to 4 ppm.
- 35 2. A non-aqueous electrolyte secondary battery of claim 1 wherein a signal strength of said NMR signals appearing within the range of -10 to 4 ppm are 1 to 10 times of a NMR signal strength appearing within the range of 10 to 20 ppm.

FIG. 1



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## INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP00/02502

A. CLASSIFICATION OF SUBJECT MATTER Int.Cl <sup>7</sup> H01M4/38, H01M4/02, H01M10/40		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) Int.Cl <sup>7</sup> H01M4/38, H01M4/02, H01M10/40		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Jitsuyo Shinan Koho 1926-1996 Toroku Jitsuyo Shinan Koho 1994-2000 Kokai Jitsuyo Shinan Koho 1971-2000 Jitsuyo Shinan Toroku Koho 1996-2000		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP, 0883199, A (Matsushita Electric Industrial Co., Ltd.), 09 December, 1998 (09.12.98), claims 1-11, etc. & JP, 2000-30703, A	1, 2
A	JP, 10-208741, A (Hitachi, Ltd.), 07 August, 1998 (07.08.98), Claims 1, 2; Par. No. 0013, etc. (Family: none)	1, 2
A	JP, 10-321225, A (Hitachi, Ltd.), 04 December, 1998 (04.12.98), Claims 1, 2, 3, 5; Par. No. 0014, etc. & US, 6030726, A & KR, 98086348, A	1, 2
A	JP, 10-316426, A (Tokuyama Corp.), 02 December, 1998 (02.12.98), Claims 1 to 4; Par. Nos. 0016 to 0036, etc. (Family: none)	1, 2
PA	JP, 11-297311, A (Kao Corporation), 29 October, 1999 (29.10.99),	1, 2
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search 05 July, 2000 (05.07.00)		Date of mailing of the international search report 18 July, 2000 (18.07.00)
Name and mailing address of the ISA/ Japanese Patent Office		Authorized officer
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International application No.

PCT/JP00/02502

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
	Claims 1 to 4, etc. (Family: none)	

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